



Nutrición Hospitalaria



Trabajo Original

Obesidad y síndrome metabólico

Visceral adiposity index associated with behavioral and inflammatory parameters in adults: a population based study

Índice de adiposidad visceral asociado con parámetros inflamatorios y del comportamiento: estudio transversal de población

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Abstract

Introduction: the visceral adiposity index (VAI) is an indicator of fat distribution and function which is considered a predictor of cardiometabolic risk.

Objective: analyze the factors associated with VAI in Brazilian adults.

Methods: a cross-sectional population-based study was conducted with 854 adults, in Viçosa, MG, Brazil. A questionnaire was applied and anthropometric measurements, body composition and biochemical data were collected. Ordinal logistic regression was used to evaluate the factors associated with VAI.

Results: the increase in percentage of fat, uric acid concentration and ultra-sensitive C reactive protein in the blood was positively associated with VAI in males. The level of physical activity in leisure was negatively associated. Among women, the increase in age, neck circumference, sitting time of more than 300 minutes, increased serum uric acid concentration and ultra-sensitive C reactive protein were associated with VAI.

Conclusion: except for age, all other factors associated with VAI are modifiable, suggesting that adopting early intervention measures that promote changes in habits and alter the risk of increased visceral adiposity and consequently the appearance of comorbidities should be encouraged.

Key words:

Adipose tissue.
Investigation.
Anthropometry.
Comorbidities.

Resumen

Introducción: el índice de adiposidad visceral (VAI) es un indicador de distribución y función de la grasa que se considera un predictor del riesgo cardiometabólico.

Objetivo: analizar los factores asociados con el VAI en adultos brasileños.

Métodos: se realizó un estudio transversal de población con 854 adultos, en Viçosa, MG, Brasil. Se aplicó un cuestionario y se analizaron las medidas antropométricas, la composición corporal y los datos bioquímicos. La regresión logística ordinal se utilizó para evaluar los factores asociados con VAI.

Resultados: el aumento en el porcentaje de grasa, la concentración de ácido úrico y la proteína C reactiva ultrasensible en la sangre se asoció positivamente con VAI en varones. El nivel de actividad física en el ocio se asoció negativamente. Entre las mujeres, el aumento de la edad, la circunferencia del cuello, el tiempo en sedestación de más de 300 minutos, el aumento de la concentración sérica de ácido úrico y la proteína C reactiva ultrasensible se asociaron con VAI.

Conclusión: con excepción de la edad, todos los otros factores asociados con el VAI son modificables, lo que sugiere que debe fomentarse la adopción de medidas de intervención temprana que promuevan cambios en los hábitos y alteren el riesgo de aumento de la adiposidad visceral y, consecuentemente, la aparición de comorbilidades.

Palabras clave:

Tejido adiposo.
Investigación.
Antropometría.
Comorbilidades.

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INTRODUCTION

The visceral fat component is the adipose tissue compartment most strongly associated with cardiometabolic complications (1-3), but its precise quantification requires high cost imaging studies, making its use in population studies and clinical practice unfeasible. For this reason, there is an increasing search for a simple, inexpensive, and easy-to-apply tool to assess visceral adiposity.

Because excess visceral fat is an independent risk factor for the development of cardiovascular diseases and is related to associated with dyslipidemia, systemic arterial hypertension, insulin resistance, and subclinical inflammation (4,5), the formulation of an indicator that can be routinely applicable is of importance to evaluate this fat as it would allow its wide use.

Recently, the visceral adiposity index (VAI) has been proposed as an indicator of fat distribution and function (6) and considered as a predictor of cardiometabolic risk (7). VAI is gender-specific and takes up simple anthropometric parameters such as waist circumference (WC) and body mass index (BMI), as well as the metabolic parameters triglyceride (TG) and high-density lipoprotein (HDL-c). VAI showed a high correlation with the visceral fat measured by magnetic resonance (6), in addition to association with cardiometabolic risk (8), metabolic syndrome (9), diabetes (10), and hypertension (11).

Population-based studies have not yet been conducted to determine which factors are associated with VAI in the Brazilian adult population and how this occurs, with his gap remaining unexplained. Therefore, the objective of this study was to analyze the association of socioeconomic, demographic, anthropometric, body composition, behavioral and biochemical factors with VAI in Brazilian adults.

MATERIALS AND METHODS

STUDY DESIGN AND SAMPLE CALCULATION

The current research was conducted as a cross-sectional, population-based study from 2012 to 2014. Data were obtained from a research that aimed to evaluate the health conditions of the adult population of Viçosa, Minas Gerais, Brazil. The study was approved by the Ethics Committee in Research of the Federal University of Viçosa (Regulation 02/2013 / CEP / 07.12.13) and its procedures are described in Segheto et al. (12).

Study sample consisted of adult individuals aged 20 to 59 years, of both sexes and living in the urban area of the municipality. For the sample calculation of this outcome, 43,431 individuals were considered as reference population (13), confidence level of 95.0%, expected prevalence of 50%, predicted sample error of 4.5 percentage points, and a design effect of 1.5. The obtained value was added with 10% of losses and refusals and 10% more to control confounding factors, requiring a minimum sample of 844 individuals. As this study is part of a survey that aims to evaluate different outcomes, 854 individuals were included who

had the complete data for the calculation of VAI and $TG \leq 279$ mg/dL and $BMI \leq 40$ kg/m² (14). The sampling process was probabilistic, without replacement, and used a two-stage conglomerate sampling (census and domicile).

DATA COLLECTION

Data collection was carried out in two stages: the first stage was conducted in the domiciles, by applying a structured questionnaire, while the second stage was carried out in the university's premises, using the anthropometric evaluations body composition and biochemical parameters. Initially, a total of 30 census tracts were selected among the 99 existing in the urban area of the municipality, then, the blocks and the starting corner of the data collection were randomly chosen. After identification of the households, all the adults residing in them were invited to participate in the study, considering as criteria for non-inclusion the pregnant women, individuals who were bedridden or unable to undergo anthropometric measurements and those unable to answer the questionnaires.

STUDY VARIABLES

VAI was considered the dependent variable and was calculated from the equations described below (6) and analyzed in tertile.

$$\text{Men: } \left(\frac{WC}{39.68 + (1.88 * BMI)} \right) * \left(\frac{TG}{1.03} \right) * \left(\frac{1.31}{HDL} \right)$$

$$\text{Women: } \left(\frac{WC}{36.58 + (1.89 * BMI)} \right) * \left(\frac{TG}{0.81} \right) * \left(\frac{1.52}{HDL} \right)$$

The socioeconomic, demographic, behavioral, anthropometric, body composition and biochemical characteristics were investigated as independent variables.

The socioeconomic and demographic variables included sex, age group in years, skin color categorized as white and non-white; schooling categorized into 0-4 years, 5-8 years, 9-11 years and ≥ 12 years, marital status categorized as with and without partner, and socioeconomic status grouped into higher (A + B), intermediate (C) and lower (D + E).

The variables related to the behavior were leisure-time physical activity (LTPA) and sedentary behavior. LTPA was evaluated by the International Physical Activity Questionnaire (IPAQ), long form, using the fourth domain. The LTPA was calculated by adding the time spent with moderate physical activities plus twice the time with vigorous activities, within the categories insufficiently active (< 150 minutes of activities in the week) or physically active (≥ 150 minutes of activities) (15). The sedentary behavior was evaluated by the sitting time (ST) calculated by the average of time (minutes) spent sitting on weekdays. TS was categorized into ≤ 300 minutes and > 300 minutes per day.

The anthropometric variables measured were BMI, WC and neck circumference (NC), all in triplicate, using the average. The criteria proposed by the World Health Organization (WHO) (16) were used for BMI. Normal-weight individuals were grouped into the normal weight category. WC and NC were measured respectively at the midpoint between the last rib and the iliac crest and below the laryngeal prominence.

The female body composition was estimated by the triceps, abdominal, and thigh skinfolds, and male body composition by triceps, pectoral and subscapular skinfolds. Skinfold data were used in sex-specific equation to calculate body density (17,18), then the fat percentage was estimated by the Siri equation (19). The skinfolds were measured using a Lange caliper (Cambridge Scientific Industries, Inc., Cambridge, MD), 1 mm precision.

The biochemical variables measured were HDL-c, TG, uric acid (UA) and ultra-sensitive C-reactive protein (us-CRP). Blood samples were collected for dosages after a 12-hour fast by venipuncture using a vacutainer system (Becton Dickinson, UK). HDL-c was measured by the enzymatic colorimetric method, TG and UA with commercial kits and us-CRP by the immunoturbidimetric assay.

STATISTICAL ANALYSIS

Data analysis was conducted using the statistical program Stata (Version 13.1, StataCorp, College Station, Texas). Considering

the complex nature of the sample, the command set "svy" was used, with sample weights assigned to the variables sex, age and schooling (13).

A descriptive analysis was carried out and association was found between the VAI and the independent variables by the ordinal logistic regression. After analyzing the independent variables in the multiple regression model, the variables that were associated with the outcome with $p < 0.05$ were maintained in the final model.

RESULTS

We evaluated 854 individuals, 392 males and 462 females. Regardless of sex, most individuals evaluated were not white (men 57.81% CI 95% 48.01-67.03 and women 60.60% CI 95% 51.83-68.73), had 12 or more years of schooling (men 55.20% CI 95% 40.11-69.38% and women 45.60% CI 95% 33.98-57.70), and were defined as intermediate socioeconomic status (men 63.42% CI95% 56.35-69.95 and women 68.44% CI95% 61.69-74.48) (Table I).

Except for LTPA in the female sex and ST in both sexes, all other parameters evaluated showed a significant difference between the third and the first VAI tertiles (Table II).

The percentage of fat, uric acid and C-reactive protein concentration in the blood were positively and independently associated

Table I. Demographic and socioeconomic characteristics of adults assessed by sex, Viçosa, MG, Brazil, 2012-2014 (n = 854)

Variable	Males		Females	
	Proportion (%)	CI (95%)	Proportion (%)	CI (95%)
Age group (years)				
20-29	39.91	(28.12-52.99)	26.65	(19.47-35.31)
30-39	26.30	(19.70-34.16)	27.24	(23.02-31.92)
40-49	16.99	(11.80-23.85)	27.28	(19.80-36.30)
50-59	16.80	(12.28-22.55)	18.83	(13.73-25.27)
Skin color				
White	42.19	(32.96-51.99)	39.40	(31.27-48.16)
Not white	57.81	(48.01-67.03)	60.60	(51.83-68.73)
Schooling (years)				
0-4	12.27	(5.91-23.73)	19.18	(11.56-30.11)
5-8	13.62	(8.56-20.99)	14.50	(10.76-19.27)
9-11	18.91	(13.41-25.98)	20.72	(17.14-24.82)
≥ 12	55.20	(40.11-69.38)	45.60	(33.98-57.70)
Marital status				
Without partner	50.03	(36.50-63.53)	48.80	(39.14-58.55)
With partner	49.97	(36.46-63.49)	51.20	(41.45-60.85)
Socioeconomic status				
Higher	31.53	(24.60-39.39)	22.51	(16.31-30.21)
Intermediate	63.42	(56.35-69.95)	68.44	(61.69-74.48)
Lower	5.05	(2.26-10.87)	9.05	(5.13-15.47)

CI: confidence interval; ≥ greater than or equal to.

Table II. Distribution (mean ± SE and 95% confidence interval) of the anthropometric variables, body composition, biochemical and behavioral variables, according to VAI tertiles in adults. Viçosa, MG, Brazil, 2012/2014 (n = 854)

Variable	Tertile 1 of VAI	Tertile 2 of VAI	Tertile 3 of VAI
<i>Age (years)</i>			
Men	31.07 ± 1.16 (28.69-33.45) ^a	34.53 ± 1.83 (30.79-38.27) ^{ab}	38.67 ± 1.68 (35.23-42.12) ^b
Women	34.16 ± 1.13 (31.84-36.49) ^a	37.43 ± 1.21 (34.94-39.91) ^{ab}	41.85 ± 1.36 (39.06-44.64) ^b
<i>NC (cm)</i>			
Men	37.18 ± 0.22 (36.73-37.61) ^a	38.53 ± 0.27 (37.98-39.08) ^b	39.08 ± 0.22 (38.64-39.53) ^b
Women	31.84 ± 0.14 (31.55-32.12) ^a	32.44 ± 0.22 (31.98-32.89) ^a	33.85 ± 0.22 (33.39-34.31) ^b
<i>% Fat</i>			
Men	15.73 ± 0.74 (14.20-17.25) ^a	20.50 ± 0.63 (19.21-21.80) ^b	23.22 ± 0.42 (22.35-24.08) ^c
Women	29.90 ± 0.58 (28.71-31.08) ^a	32.01 ± 0.54 (30.90-33.12) ^{ab}	34.23 ± 0.58 (33.04-35.43) ^b
<i>UA (mg/dL)</i>			
Men	4.35 ± 0.12 (4.11-4.60) ^a	4.52 ± 0.11 (4.30-4.74) ^a	4.99 ± 0.09 (4.80-5.18) ^b
Women	3.03 ± 0.08 (2.87-3.19) ^a	3.18 ± 0.07 (3.04-3.33) ^a	3.79 ± 0.16 (3.48-4.12) ^b
<i>us-CRP (mg/L)</i>			
Men	1.05 ± 0.13 (0.79-1.31) ^a	1.32 ± 0.14 (1.02-1.61) ^{ab}	1.64 ± 0.13 (1.37-1.91) ^b
Women	1.58 ± 0.16 (1.25-1.91) ^a	1.82 ± 0.16 (1.49-2.14) ^a	2.58 ± 0.16 (2.25-2.92) ^b
<i>LTPA (min)</i>			
Men	194.90 ± 30.29(132.85-256.96) ^a	175.05 ± 39.06 (95.03-255.07) ^{ab}	82.71 ± 19.73 (42.29-123.13) ^b
Women	75.33 ± 15.88 (42.79-107.87)	113.55 ± 21.26 (70.00-157.10)	69.90 ± 15.95 (37.23-102.56)
<i>ST (min)</i>			
Men	344.35 ± 33.68 (275.36-413.33)	334.88 ± 23.68 (286.36-383.39)	329.31 ± 12.07 (304.59-354.04)
Women	299.06 ± 20.27 (257.53-340.59)	274.39 ± 23.27 (226.52-322.26)	277.58 ± 23.07 (230.32-324.85)

SE: standard error; VAI: visceral adiposity index - VAI Tertile of men (tertile 1: 0.211-0.889; tertile 2: 0.891-1.667; tertile 3: 1.676-7.840) - VAI Tertile of women (tertile 1: 0.309-1.049; tertile 2: 1.059-1.667; tertile 3: 1.673-1.750); NC: neck circumference; %: Percentage; UA: uric acid; us-CRP: ultra-sensitive c-reactive protein; LTPA: leisure-time physical activity; ST: sitting time. Bold numbers and different letters indicate statistical difference.

with VAI in the male sex, while the leisure-time physical activity level was independent and negatively associated with VAI. A positive and independent association was found in females for increase in age, neck circumference, sitting time over 300 minutes, increased uric acid concentration and C-reactive protein in blood (Table III).

DISCUSSION

This is the first study evaluating the association of socioeconomic, demographic, anthropometric, body composition, behavioral and biochemical factors with VAI in Brazilian adults and may serve as a basis for conducting longitudinal studies aimed to assess the risk of increase in VAI and its association with comorbidities.

We found a positive and independent association between VAI and increase in age for women (OR 1.04 CI 95% 1.01-1.07), with a significant increase in age between the third (41.85 ± 1.361 CI 95% 39.06-44.64) and the first tertiles (34.16 ± 1.135 95% CI 31.84-36.49) of VAI in both sexes.

Previous studies (20,21) have shown that advancing age leads to an increase in visceral adiposity, reinforcing the need to evaluate this adiposity in different age groups. Possible explanations for this association in women are changes in lifestyle that promote decreased levels of physical activity and consequently change in basal metabolism, as well as changes in hormone levels in particular after menopause (21).

The increase in the percentage of total fat was positively associated in men (OR 1.14 95% CI 1.09-1.18), indicating that an increase in overall adiposity in this group also leads to an increase in visceral adiposity, which may compromise health (22,23).

We found that increase in NC was also positively associated with VAI independently in women (OR 1.28 95% CI 1.15-1.44). NC is a simple marker of upper body adiposity, and a synergistic effect between this measure and visceral adipose tissue with cardiometabolic risk factors was observed in the Framingham Heart Study (24). It is worth noting that fat located in the upper part of the body can cause metabolic abnormalities, increasing circulating free fatty acids, which can lead to insulin resistance (25,26).

Table III. Final model of the ordinal logistic regression for factors associated with the visceral adiposity index in adults. Viçosa, MG, Brazil, 2012-2014 (n = 854)

Variable	Males			Females		
	OR	CI 95%	p	OR	CI 95%	p
Age (years)	—	—		1.04	1.01-1.07	0.003
% fat	1.14	1.09-1.18	< 0.001	—	—	
NC (cm)	—	—		1.28	1.15-1.44	< 0.001
LTPA (min)			0.007			
Inactive	1.00	—		—	—	
Active	0.54	0.35-0.83		—	—	
ST (min)						0.023
≤ 300	—	—		1.00	—	
> 300	—	—		2.10	1.11-3.97	
UA (mg/dL)	1.42	1.14-1.78	0.003	1.50	1.14-1.98	0.006
us-CRP (mg/L)	1.19	1.05-1.35	0.008	1.19	1.06-1.33	0.005

OR: odds ratio; CI: confidence interval; %: Percentage; NC: neck circumference; LTPA: leisure-time physical activity; ST: sitting time; UA: uric acid; us-CRP: ultra-sensitive C-reactive protein.

In our study, we found that being active contributed to a 46% reduction in the chance of increased visceral adiposity among men (OR: 0.54 95% CI 0.35-0.83), constituting a protective factor. LTPA was statistically lower in men who had higher VAIs (82.71 ± 19.73 CI 95% 42.29-123.13) than in those with lower VAI (194.90 ± 30.29 CI 95% 132.85-256.96). Inverse association between LTPA and obesity has previously been demonstrated (27), reinforcing the benefits of an active life in body composition.

Our results also showed that sedentary behavior time > 300 minutes in women was independently and positively associated with VAI (OR 2.10 95% CI 1.11-3.97). Previous studies have shown association between greater time spent in sedentary behavior and greater waist circumference (1.89 95% CI 0.94-2.83 cm) (28), as well as higher risk of overweight and obesity (OR 1.38 95% CI 1.26-1.52) (29). The physiological effect of sedentary behavior promoting unhealthiness may result from the absence of muscle contraction that leads to an increase in triglycerides, plasma glucose and a reduction in lipoprotein lipase activity, which is an enzyme related to the regulation of triglyceride absorption and the production of HDL-c in skeletal muscles (30).

Identifying differences between the sexes in the association of LTPA and sedentary behavior with VAI is important for proposing specific intervention strategies for this adult population. This study suggests that physical activity should be encouraged for men to reduce the chance of increasing VAI, while for women the greatest encouragement should be to decrease sitting time as this more than doubles the chance of increasing VAI.

We observed that in both sexes the increase in UA concentration was positively associated (OR: 1.42 CI 95% 1.14-1.78 in men; OR: 1.50 CI 95% 1.14-1.98 in women). The association between high concentration of UA with factors such as obesity, hypertriglyceridemia, hypertension, diabetes and consequently metabolic syndrome has been demonstrated in the Brazilian adult population (31), as well as a correlation between VAI and UA (9)

in adult Chinese, suggesting the possibility of using VAI to predict inflammation. Recently, an association between increased UA and visceral and subcutaneous adipose tissue was described, but the association was lost when the effect of visceral adipose tissue was controlled (32).

Increase in us-CRP concentration was also positively associated in both sexes (OR: 1.19 95% CI 1.05-1.35 in men; OR: 1.19 CI 95% 1.06-1.33 in women). As us-CRP is a marker for low-grade inflammation, its association with IAV suggests that it may also indicate inflammation. Recent research on the impact of the number of risk factors for metabolic syndrome on the serum concentration of us-CRP in women and men has shown that even in absence of evident disease and pharmacological treatment, individuals with risk factors for metabolic syndrome already have an increased concentration of us-CRP (33). Correlation between VAI and us-CRP in women (0.19 $p < 0.005$) has been previously demonstrated (34), corroborating with our study, as well as the increase observed in us-CRP as the tertile value of VAI increases both in pre-diabetes (OR 2,17 CI 95% 1,40-3,37) and type 2 diabetes (OR 1,17 95% CI 1,16-2,71) (35).

Some limitations of this study are that the transversal design does not allow us to establish a temporal relation between the exposure and the outcome, but this was not the objective of our study. The proportion of selected individuals had no correspondence to data referring to the Viçosa population for sex, schooling, and age, which was corrected by weighing the sample means. Finally, a questionnaire was used to estimate LTPA and quantification of sitting time as an indicator of sedentary behavior was memory dependent. However, we believe that when we evaluated LTPA with a validated questionnaire and took only the last week for the sitting time we were able to minimize these limitations.

Considering the data from this study, we conclude that there is a positive and independent association in men between VAI and the increase in fat percentage, in the concentration of UA

and us-CRP in the blood, whereas a negative association was observed with LTPA. In women, the increase in age and NC, sitting time greater than 300 minutes, increase in AU concentration and us-CRP were the independent factors and positively associated with VAI. With the exception of age, the other factors are modifiable, suggesting that the adoption of early intervention measures that lead to changes in habits may alter the risk of increased visceral adiposity and, consequently, the appearance of comorbidities. However, more research on this topic needs to be undertaken to investigate the factors associated with VAI in different populations.

REFERENCES

1. Pinho CPS. Tecido adiposo visceral e subcutâneo em adultos com excesso de peso: aspectos metodológicos, metabólicos e terapêuticos. Universidade Federal de Pernambuco; 2016.
2. Pak K, Lee S, Lee J, Seok J, Kim I. Comparison of Visceral Fat Measures with Cardiometabolic Risk Factors in Healthy Adults. *PLoS One* 2016;11:e0153031.
3. Shah RV, Murthy VL, Abbasi SA, Blankstein R, Kwong RY, Goldfine AB, et al. Visceral Adiposity and the Risk of Metabolic Syndrome Across Body Mass Index. *JACC Cardiovasc Imaging* 2014;7:1221-35. DOI: 10.1016/j.jcmg.2014.07.017
4. Lemieux I, Poirier P, Bergeron J, Alméras N, Lamarche B, Cantin B, et al. Hypertriglyceridemic waist: a useful screening phenotype in preventive cardiology? *Can J Cardiol* 2007;23:23-31. DOI: 10.1016/S0828-282X(07)71007-3
5. Després J, Lemieux I, Bergeron J, Pibarot P, Mathieu P, Larose E, et al. Abdominal Obesity and the Metabolic Syndrome : Contribution to Global Cardiometabolic Risk. *Arterioscler Thromb Vasc Biol* 2008;28:1039-49. DOI: 10.1161/ATVBAHA.107.159228
6. Amato MC, Giordano C, Galia M, Criscimanna A, Vittabile S, Midiri M, et al. Visceral Adiposity Index: a reliable indicator of visceral fat function associated with cardiometabolic risk. *Diabetes Care* 2010;33:920-2. DOI: 10.2337/dc09-1825
7. Amato MC, Giordano C. Visceral adiposity index: An indicator of adipose tissue dysfunction. *Int J Endocrinol* 2014;1-7. DOI: 10.1155/2014/730827
8. Ahmad MN, Haddad FH. Suitability of visceral adiposity index as a marker for cardiometabolic risks in Jordanian adults. *Nutr Hosp* 2015;32:2701-9. DOI: 10.3305/nh.2015.32.6.9543
9. Chen G-P, Qi J-C, Wang B-Y, Lin X, Zhang X-B, Zhao J-M, et al. Applicability of visceral adiposity index in predicting metabolic syndrome in adults with obstructive sleep apnea: a cross-sectional study. *BMC Pulm Med* 2016;16. DOI: 10.1186/s12890-016-0198-0
10. Chen C, Xu Y, Guo ZR, Yang J, Wu M, Hu XS. The application of visceral adiposity index in identifying type 2 diabetes risks based on a prospective cohort in China. *Lipids Health Dis* 2014;13. DOI: 10.1186/1476-511X-13-108
11. Ding Y, Gu D, Zhang Y, Han W, Liu H, Qu Q. Significantly Increased Visceral Adiposity Index in Prehypertension. *PLoS One* 2015;10:e0123414. DOI: 10.1371/journal.pone.0123414
12. Segheto W, Silva DCG, Coelho FA, Reis VG, Morais SHO, Marins JCB, et al. Body adiposity index and associated factors in adults: method and logistics of a population-based study. *Nutr Hosp* 2015;32:101-9. DOI: 10.3305/nh.2015.32.1.8391
13. Instituto Brasileiro de Geografia e Estatística (IBGE). Censo Demográfico 2010. Características da população e dos domicílios; 2010.
14. Amato MC, Giordano C. Clinical indications and proper use of Visceral Adiposity Index. *Nutr Metab Cardiovasc Dis* 2013;23:e31-2. DOI: 10.1016/j.numecd.2013.04.006
15. Haskell WL, Lee I-M, Pate RR, Powell KE, Blair SN, Franklin BA, et al. Physical Activity and Public Health Updated Recommendation for Adults From the American College of Sports Medicine and the American Heart Association. *Med Sci Sport Exerc* 2007;39:1423-34. DOI: 10.1161/CIRCULATIONAHA.107.185649
16. World Health Organization. Obesity: preventing and managing the global epidemic. Report of a WHO consultation. (WHO Technical Report Series 894) 2000;894:i-xii, 1-253. DOI: 10.1016/S0140-6736(57)91352-1
17. Jackson AS, Pollock ML, Ward A. Generalized equations for predicting body density of women. *Med Sci Sports Exerc* 1980;12:175-81.
18. Jackson AS, Pollock ML. Generalized equations for predicting body density of men. *Br J Nutr* 1978;40:497-504.
19. Siri WE. The gross composition of the body. *Adv Biol Med Phys* 1956;4: 239-80.
20. Kotani K, Tokunaga K, Fujioka S, Kobatake T, Keno Y, Yoshida S, et al. Sexual dimorphism of age-related changes in whole-body fat distribution in the obese. *Int J Obes Relat Metab Disord* 1994;18:202-7.
21. Shen W, Punyanitya M, Silva AM, Chen J, Gallagher D, Sardinha LB, et al. Sexual dimorphism of adipose tissue distribution across the lifespan: a cross-sectional whole-body magnetic resonance imaging study. *Nutr Metab (Lond)* 2009;6:17. DOI: 10.1186/1743-7075-6-17
22. Patel P, Abate N. Body Fat Distribution and Insulin Resistance. *Nutrients* 2013;5:2019-27. DOI: 10.3390/nu5062019
23. Després J-P. Body Fat Distribution and Risk of Cardiovascular Disease: An Update. *Circulation* 2012; 126:1301-13. DOI: 10.1161/CIRCULATIONAHA.111.067264
24. Preis SR, Massaro JM, Hoffmann U, D'Agostino RB, Levy D, Robins SJ, et al. Neck Circumference as a Novel Measure of Cardiometabolic Risk: The Framingham Heart Study. *J Clin Endocrinol Metab* 2010;95:3701-10. DOI: 10.1210/jc.2009-1779
25. Nielsen S, Guo Z, Johnson CM, Hensrud DD, Jensen MD. Splanchnic lipolysis in human obesity. *J Clin Invest* 2004;113:1582-8. DOI: 10.1172/JCI21047
26. Lee J, Pedley A, Therkelsen K, Hoffmann U, Massaro J, Levy D, et al. Upper body subcutaneous fat is associated with cardiometabolic risk factors. *Am J Med* 2017;130(8):958-966.e1. DOI: 10.1016/j.amjmed.2017.01.044
27. Banks E, Lim L, Seubsman S-A, Bain C, Sleight A. Relationship of obesity to physical activity, domestic activities, and sedentary behaviours: cross-sectional findings from a national cohort of over 70,000 Thai adults. *BMC Public Health* 2011;11:762. DOI: 10.1186/1471-2458-11-762
28. Cooper AR, Sebire S, Montgomery AA, Peters TJ, Sharp DJ, Jackson N, et al. Sedentary time, breaks in sedentary time and metabolic variables in people with newly diagnosed type 2 diabetes. *Diabetologia* 2012;55:589-99. DOI: 10.1007/s00125-011-2408-x
29. Duncan MJ, Vandelanotte C, Caperchione C, Hanley C, Mummery WK. Temporal trends in and relationships between screen time, physical activity, overweight and obesity. *BMC Public Health* 2012;12:1060. DOI: 10.1186/1471-2458-12-1060
30. Bey L, Hamilton MT. Suppression of skeletal muscle lipoprotein lipase activity during physical inactivity: a molecular reason to maintain daily low-intensity activity. *J Physiol* 2003;551:673-82. DOI: 10.1113/jphysiol.2003.045591
31. Rodrigues SL, Baldo MP, Capingana P, Magalhães P, Dantas EM, Molina M del CB, et al. Distribuição por gênero de ácido úrico sérico e fatores de risco cardiovascular: estudo populacional. *Arq Bras Cardiol* 2012;98:13-21. DOI: 10.1590/S0066-782X2011005000116
32. Bertoli S, Leone A, Vignati L, Spadafranca A, Bedogni G, Vanzulli A, et al. Metabolic correlates of subcutaneous and visceral abdominal fat measured by ultrasonography: a comparison with waist circumference. *Nutr J* 2016;15:2. DOI: 10.1186/s12937-015-0120-2
33. Garcia VP, Rocha HNM, Sales ARK, Rocha NG, da Nóbrega ACL. Sex Differences in High Sensitivity C-Reactive Protein in Subjects with Risk Factors of Metabolic Syndrome. *Arq Bras Cardiol* 2016;106:182-7. DOI: 10.5935/abc.20160027
34. Al-Daghri NM, Al-Attas OS, Alokail MS, Alkharfy KM, Charalampidis P, Livadadas S, et al. Visceral adiposity index is highly associated with adiponectin values and glycaemic disturbances. *Eur J Clin Invest* 2013;43:183-9. DOI: 10.1111/eci.12030
35. Liu PJ, Ma F, Lou HP, Chen Y. Visceral Adiposity Index Is Associated with Pre-Diabetes and Type 2 Diabetes Mellitus in Chinese Adults Aged 20-50. *Ann Nutr Metab* 2016;68:235-43. DOI: 10.1159/000446121